AMENDMENTS TO THE CLAIMS

This listing of claims will replace all prior versions and listings of claims in the application:

LISTING OF CLAIMS:

1. (currently amended): A compound of formula (I):

wherein

ring A, ring B, and ring D each independently represents a cyclic group which may be substituted;

J represents a bond or a spacer having 1 to 8 atoms in its main chain; and

G represents a bond or a spacer having 1 to 4 atoms in its main chain;

wherein

<u>is</u>

$$\mathsf{R}^\mathsf{D} \biguplus_{\mathsf{N}}^\mathsf{N} \bigvee_{\mathsf{G}}^\mathsf{J} \qquad \text{or} \qquad \biguplus_{\mathsf{N}}^\mathsf{N} \bigvee_{\mathsf{G}}^\mathsf{J}$$

wherein R^D represents a substituent of ring D;

represents a single bond or a double bond; and

M represents a 3- to 11-membered monocyclic or bicyclic cyclic group which may be substituted;

ring B is a C₃₋₈ monocyclic carbocyclic ring which may be substituted or a 3- to 8-membered monocyclic heterocyclic ring having 1 to 4 nitrogen atoms, 1 or 2 oxygen atoms and/or 1 or 2 sulfur atoms as a hetero atom(s) which may be substituted;

J is

$$-0$$
 $\stackrel{R^3}{\stackrel{}{\checkmark}}$ R^4

wherein R³ and R⁴ each independently represents hydrogen or C₁₋₈ alkyl; and

E represents a bond or a spacer having 1 to 6 atoms in its main chain;

G is -NR^{T1}-SO₂-

wherein R^{T1} represents hydrogen, C_{1-8} alkyl which may be substituted, C_{2-8} alkenyl which may be substituted, C_{2-8} alkynyl which may be substituted or a 3- to 8-membered cyclic group which may be substituted;

or a salt thereof.

Claims 2-6. (canceled).

7. (currently amended): The compound according to claim 61, wherein

is

$$R^{D}$$
 N
 G
 R^{D}
 N
 G
 N

wherein

R^D has the same meaning as described in claim 61.

- 8. (original): The compound according to claim 1, wherein ring A is a carbocyclic ring which may be substituted.
- 9. (original): The compound according to claim 1, wherein ring A is a heterocyclic ring which may be substituted.
- 10. (original): The compound according to claim 8, wherein the carbocyclic ring is a C_{3-15} monocyclic, bicyclic or tricyclic carbocyclic ring.
- 11. (original): The compound according to claim 9, wherein the heterocyclic ring is a 3-to 15-membered monocyclic, bicyclic or tricyclic heterocyclic ring having 1 to 4 nitrogen atoms, 1 or 2 oxygen atoms and/or 1 or 2 sulfur atoms as a hetero atom(s).
- 12. (original): The compound according to claim 10, wherein the carbocyclic ring is a benzene ring or a naphthalene ring.

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13. (original): The compound according to claim 11 wherein the heterocyclic ring is a

pyridine ring, a pyrazole ring, a dioxaindane ring or a benzodioxane ring.

Claims 14 - 19. (canceled).

20. (currently amended): The compound according to claim 181, wherein the

C₃₋₈ monocyclic carbocyclic ring represented by ring B is a benzene ring.

21. (currently amended): The compound according to claim 191, wherein the 3- to 8-

membered monocyclic heterocyclic ring having 1 to 4 nitrogen atoms, 1 or 2 oxygen atoms

and/or 1 or 2 sulfur atoms as a hetero atom(s) represented by ring B is a pyridine ring or a

thiophene ring.

Claims 22-24. (canceled).

25. (currently amended): The compound according to claim 241, wherein R³ and R⁴

each independently represents hydrogen or methyl.

26. (currently amended): The compound according to claim 241, wherein E is a bond[[,]].

27. (currently amended): The compound according to claim 241, wherein E is a spacer

having 1 to 6 atoms in its main chain.

28. (original): The compound according to claim 27, wherein E is C₁₋₄ alkylene or

 C_{1-3} alkyleneoxy.

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- 29. (original): The compound according to claim 28, wherein E is methylene or methylenoxy.
 - 30. (canceled).
 - 31. (canceled).
- 32. (currently amended): The compound according to claim 311, wherein G is -NH-SO₂-.
- 33. (currently amended): The compound according to claim 1, wherein the compound is a compound of formula (A):

wherein

 R^1 and R^2 each independently represents (1) hydrogen, (2) C_{1-8} alkyl, (3) C_{2-8} alkenyl, (4) C_{2-8} alkynyl, (5) halogen, (6) cyano, (7) nitro, (8) -CONR⁷R⁸, (9) -COOR⁹, (10) Cyc1 or (11) C_{1-8} alkyl substituted with 1 to 5 groups selected from (a) -CONR⁷R⁸, (b) -COOR⁹, (c) -OR¹⁰, (d) -NR¹¹R¹², (e) halogen, and (f) Cyc1; or

 R^1 and R^2 are taken together to represent $C_{3.4}$ alkylene, -CH=CH-CH₂-, -CH₂-CH=CH-, -CH=CH-CH=CH- or -CH=CH-CH₂-CH₂-, wherein the carbocyclic ring to be formed may be substituted with $C_{1.8}$ alkyl, $C_{2.8}$ alkenyl, $C_{2.8}$ alkynyl, $C_{1.8}$ alkoxy, halogen, cyano, nitro or

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hydroxyl, wherein R⁷ and R⁸ each independently represents (1) hydrogen, (2) C₁₋₈ alkyl, (3) C₂₋₈ alkenyl, (4) C₂₋₈ alkynyl, (5) Cyc2, (6) -OR¹³ or (7) C₁₋₈ alkyl, C₂₋₈ alkenyl or C₂₋₈ alkynyl substituted with 1 to 5 groups selected from (a) -OR¹³, (b) -NR¹⁴R¹⁵, (c) -NR¹⁶COR¹⁷, (d) halogen, (e) CF₃, and (f) Cyc2; or R⁷ and R⁸ are taken together with the adjacent nitrogen atom to represent a 3- to 8-membered monocyclic heterocyclic ring having at least one nitrogen atom as a hetero atom(s) and 0 to 3 nitrogen atoms, 0 to 1 oxygen atom and/or 0 to 1 sulfur atom as an other hetero atom(s), wherein the heterocyclic ring may be substituted with (a) C₁₋₈ alkyl, (b) halogen, (c) hydroxyl, or (d) C₁₋₈ alkyl substituted with hydroxyl;

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 R^{13} to R^{17} each independently represents (1) hydrogen, (2) C_{1-8} alkyl, (3) C_{2-8} alkenyl, (4) C_{2-8} alkynyl, (5) Cyc1, or (6) C_{1-8} alkyl, C_{2-8} alkenyl or C_{2-8} alkynyl substituted with Cyc1;

 R^9 to R^{12} each independently represents (1) hydrogen, (2) C_{1-8} alkyl, (3) C_{2-8} alkenyl, (4) C_{2-8} alkynyl, (5) Cyc1, or (6) C_{1-8} alkyl, C_{2-8} alkenyl or C_{2-8} alkynyl substituted with Cyc1;

Cyc1 represents a C_{3-15} monocyclic, bicyclic or tricyclic carbocyclic ring or a 3- to 15-membered monocyclic, bicyclic or tricyclic heterocyclic ring having 1 to 4 nitrogen atoms, 1 or 2 oxygen atoms and/or 1 or 2 sulfur atoms as a hetero atom(s), wherein Cyc1 may be substituted with 1 to 5 of \mathbb{R}^{18} ;

 R^{18} represents (1) C_{1-8} alkyl, (2) C_{2-8} alkenyl, (3) C_{2-8} alkynyl, (4) halogen, (5) cyano, (6) nitro, (7) trifluoromethyl, (8) trifluoromethoxy, (9) $-OR^{19}$, (10) $-SR^{20}$, (11) $-NR^{21}R^{22}$, (12) $-COR^{23}$, (13) $-COOR^{24}$, (14) $-NR^{25}COR^{26}$, (15) $-CONR^{27}R^{28}$, (16) Cyc2, or (17) C_{1-8} alkyl, C_{2-8} alkenyl or C_{2-8} alkynyl substituted with 1 to 5 groups selected from (a) halogen, (b) cyano, (c) nitro, (d) trifluoromethyl, (e) trifluoromethoxy, (f) $-OR^{19}$, (g) $-SR^{20}$, (h) $-NR^{21}R^{22}$, (i) $-COR^{23}$, (j) $-COOR^{24}$, (k) $-NR^{25}COR^{26}$, (l) $-CONR^{27}R^{28}$, and (m) Cyc2;

 R^{19} to R^{28} each independently represents (1) hydrogen, (2) C_{1-8} alkyl, (3) C_{2-8} alkenyl, (4) C_{2-8} alkynyl, (5) Cyc2, or (6) C_{1-8} alkyl, C_{2-8} alkenyl or C_{2-8} alkynyl substituted with Cyc2;

Cyc2 represents a C_{3-8} monocyclic carbocyclic ring or a 3- to 8-membered monocyclic heterocyclic ring having 1 to 4 nitrogen atoms, 1 or 2 oxygen atoms and/or 1 or 2 sulfur atoms as a hetero atom(s), wherein Cyc2 may be substituted with 1 to 5 of R^{29} ;

 R^{29} represents (1) C_{1-8} alkyl, (2) C_{2-8} alkenyl, (3) C_{2-8} alkynyl, (4) halogen, (5) cyano, (6) nitro, (7) hydroxyl, (8) trifluoromethyl, (9) trifluoromethoxy, or (10) -OR¹⁰⁰;

 R^{100} represents C_{1-8} alkyl.;

R³ and R⁴ each independently represents hydrogen or C₁₋₈ alkyl;

 E^1 represents a bond or C_{1-6} alkylene, wherein a carbon atom in the alkylene group may be substituted with oxygen, sulfur, or -NR³⁰-;

 R^{30} represents (1) C_{1-8} alkyl, (2) C_{2-8} alkenyl, (3) C_{2-8} alkynyl, (4) phenyl, or (5) C_{1-8} alkyl substituted with phenyl;

ring A^1 represents a C_{3-15} monocyclic, bicyclic or tricyclic carbocyclic ring or a 3- to 15-membered monocyclic, bicyclic or tricyclic heterocyclic ring having 1 to 4 nitrogen atoms, 1 or 2 oxygen atoms and/or 1 or 2 sulfur atoms as a hetero atom(s);

 R^5 represents (1) C_{1-8} alkyl, (2) C_{2-8} alkenyl, (3) C_{2-8} alkynyl, (4) halogen, (5) cyano, (6) nitro, (7) trifluoromethyl, (8) trifluoromethoxy, (9) $-OR^{31}$, (10) $-NR^{32}R^{33}$, (11) $-NR^{34}COR^{35}$, (12) Cyc3, or (13) C_{1-8} alkyl, C_{2-8} alkenyl or C_{2-8} alkynyl substituted with 1 to 5 groups selected from (a) halogen, (b) cyano, (c) nitro, (d) trifluoromethyl, (e) trifluoromethoxy, (f) $-OR^{31}$, (g) $-NR^{32}COR^{33}$, (h) $-NR^{34}COR^{35}$, and (i) Cyc3;

 R^{31} to R^{35} each independently represents (1) hydrogen, (2) C_{1-8} alkyl, (3) C_{2-8} alkenyl, (4) C_{2-8} alkynyl, (5) Cyc3, or (6) C_{1-8} alkyl, C_{2-8} alkenyl or C_{2-8} alkynyl substituted with 1 to 5 groups selected from (a) Cyc3, (b) -OR 36 and (c) -NR 37 R 38 ;

 R^{36} to R^{38} each independently represents (1) hydrogen, (2) C_{1-8} alkyl, (3) -OR³⁹, or (4) -NR⁴⁰R⁴¹:

R³⁹ to R⁴¹ each independently represents hydrogen or C₁₋₈ alkyl;

Cyc3 represents a C₃₋₈ monocyclic carbocyclic ring or a 3- to 8-membered monocyclic heterocyclic ring having 1 to 4 nitrogen atoms, 1 or 2 oxygen atoms and/or 1 or 2 sulfur atoms as a hetero atom(s);

ring B^1 represents a C_{3-8} monocyclic carbocyclic ring or a 3- to 8-membered monocyclic heterocyclic ring having 1 to 4 nitrogen atoms, 1 or 2 oxygen atoms and/or 1 or 2 sulfur atoms as

a hetero atom(s)C₃₋₁₅ monocyclic, bicyclic or tricyclic carbocyclic ring or a 3- to 15 membered monocyclic, bicyclic or tricyclic heterocyclic ring having 1 to 4 nitrogen atoms, 1 or 2 oxygen atoms and/or 1 or 2 sulfur atoms as a hetero atom(s);

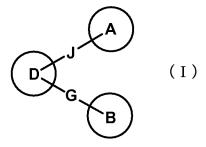
 R^6 represents (1) C_{1-8} alkyl, (2) C_{2-8} alkenyl, (3) C_{2-8} alkynyl, (4) halogen, (5) cyano, (6) nitro, (7) trifluoromethyl, (8) trifluoromethoxy, (9) $-OR^{42}$, (10) $-NR^{43}R^{44}$, (11) $-SR^{101}$, (12) $-SO_2R^{102}$, (13) $-COR^{103}$, (14) $-COOR^{104}$, (15) Cyc2, or (16) C_{1-8} alkyl, C_{2-8} alkenyl or C_{2-8} alkynyl substituted with 1 to 5 groups selected from (a) $-COOR^{104}$, (b) $-NR^{105}COR^{106}$, and (c) Cyc2;

 R^{42} to R^{44} and R^{101} to R^{106} each independently represents (1) hydrogen, (2) C_{1-8} alkyl, (3) Cyc2, or (4) -COR¹⁰⁷, or (5) C_{1-8} alkyl substituted with 1 to 5 halogen atoms;

$$R^{107}$$
 represents C_{1-8} alkyl; and

p and q each independently represents 0 or an integer of 1 to 5.

- 34. (withdrawn): A prodrug for the compound according to claim 1.
- 35. (currently amended): A pharmaceutical composition which comprises the compound of formula (I):



wherein

ring A, ring B, and ring D each independently represents a cyclic group which may be substituted;

J represents a bond or a spacer having 1 to 8 atoms in its main chain; and

G represents a bond or a spacer having 1 to 4 atoms in its main chain;

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wherein

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<u>is</u>

$$\mathsf{R}^\mathsf{D} \biguplus_{\mathsf{N}}^\mathsf{N} \bigvee_{\mathsf{G}}^\mathsf{J} \qquad \text{or} \qquad \bigvee_{\mathsf{N}}^\mathsf{N} \bigvee_{\mathsf{G}}^\mathsf{J}$$

wherein R^D represents a substituent of ring D;

represents a single bond or a double bond; and

M represents a 3- to 11-membered monocyclic or bicyclic cyclic group which may be substituted;

ring B is a C₃₋₈ monocyclic carbocyclic ring which may be substituted or a 3- to 8-membered monocyclic heterocyclic ring having 1 to 4 nitrogen atoms, 1 or 2 oxygen atoms and/or 1 or 2 sulfur atoms as a hetero atom(s) which may be substituted;

J is

$$R^3$$
 R^4 E

wherein R³ and R⁴ each independently represents hydrogen or C₁₋₈ alkyl; and

E represents a bond or a spacer having 1 to 6 atoms in its main chain;

G is -NR^{T1}-SO₂-

wherein R^{T1} represents hydrogen, C_{1-8} alkyl which may be substituted, C_{2-8} alkenyl which may be substituted, C_{2-8} alkynyl which may be substituted or a 3- to 8-membered cyclic group which may be substituted;

or a salt thereof and a pharmaceutically acceptable carrier.

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36. (currently amended): The pharmaceutical composition according to claim 35, which

is has an activity of a chemokine receptor antagonist.

37. (original): The pharmaceutical composition according to claim 36, wherein the

chemokine receptor is CCR4.

38. (currently amended): The pharmaceutical composition according to claim 37, which

is has an activity for a preventive and/or therapeutic agent for treating CCR4-mediated diseases.

39. (original): The pharmaceutical composition according to claim 38, wherein the

CCR4-mediated diseases are inflammatory and/or allergic diseases, metabolism and/or endocrine

system diseases, cancer diseases or infections.

40. (original): The pharmaceutical composition according to claim 39, wherein the

CCR4-mediated diseases are inflammatory and/or allergic diseases.

41. (original): The pharmaceutical composition according to claim 40, wherein the

inflammatory and/or allergic diseases are respiratory diseases or dermatosis.

42. (original): The pharmaceutical composition according to claim 41, wherein the

respiratory diseases are asthma.

43. (original): The pharmaceutical composition according to claim 41, wherein the

dermatosis is atopic dermatitis.

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44. (withdrawn-currently amended): A method for preventing and/or treating CCR4-

mediated diseases in a mammal, which comprises administering to a mammal an effective

amount of the compound according to claim 1 or a salt thereof.

45. (canceled).

46. (currently amended): A pharmaceutical composition which comprises: a preventive

and/or-therapeutic agent for CCR4-mediated diseases, which comprises the compound according

to claim 1 or a salt thereof as an active ingredient; and one or at least two medicaments selected

from a bronchodilator drug, a steroid drug, a non-steroidal antiinflammatory drug, a leukotriene

receptor antagonist, a phosphodiesterase inhibitor, an immunosuppressant, an anti-allergic drug,

a mediator-release inhibitor, an antihistamine drug, a metabolism promoter and/or a chemokine

inhibitor.

47. (currently amended): The pharmaceutical composition according to claim 35, which

is has an activity of inhibiting an inhibitor of effector cell function.

48. (currently amended): The pharmaceutical composition according to claim 47, which

is an inhibitor of has an activity of inhibiting cell migration function.

49. (currently amended): The pharmaceutical composition according to claim 35, which

is has an activity of regulating a TNF α regulator.

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